cholesterol, potassium, other biomarkers, etc. The biomarker concentrations in the tear film can be systematically different than the corresponding concentrations of the biomarkers in the blood, but a relationship between the two concentration levels can be established to map tear film biomarker concentration values to blood concentration levels. For example, the tear film concentration of glucose can be established (e.g., empirically determined) to be approximately one tenth the corresponding blood glucose concentration. Thus, measuring tear film analyte concentration levels provides a non-invasive technique for monitoring biomarker levels in comparison to blood sampling techniques performed by lancing a volume of blood to be analyzed outside a person's body. Moreover, the ophthalmic analyte bio-sensor platform disclosed here can be operated substantially continuously to enable real time monitoring of analyte concentrations.

[0054] To perform a reading with the system **100** configured as a tear film analyte monitor, the external reader **180** can emit radio frequency radiation **171** that is harvested to power the eye-mountable device **110** via the power supply **140**. Radio frequency electrical signals captured by the energy harvesting antenna **142** (and/or the communication antenna **170**) are rectified and/or regulated in the rectifier/regulator **146** and a regulated DC supply voltage **147** is provided to the controller **150**. The radio frequency radiation **171** thus turns on the electronic components within the eye-mountable device **110**. Once turned on, the controller **150** operates the analyte bio-sensor **162** to measure an analyte concentration level. For example, the sensor interface module **152** can apply a voltage between a working electrode and a reference electrode in the analyte bio-sensor **162**. The applied voltage can be sufficient to cause the analyte to undergo an electrochemical reaction at the working electrode and thereby generate an amperometric current that can be measured through the working electrode. The measured amperometric current can provide the sensor reading ("result") indicative of the analyte concentration. The controller **150** can operate the antenna **170** to communicate the sensor reading back to the external reader **180** (e.g., via the communication circuit **156**). The sensor reading can be communicated by, for example, modulating an impedance of the communication antenna **170** such that the modulation in impedance is detected by the external reader **180**. The modulation in antenna impedance can be detected by, for example, backscatter radiation from the antenna **170**.

[0055] In some embodiments, the system **100** can operate to non-continuously ("intermittently") supply energy to the eye-mountable device **110** to power the controller **150** and electronics **160**. For example, radio frequency radiation **171** can be supplied to power the eye-mountable device **110** long enough to carry out a tear film analyte concentration measurement and communicate the results. For example, the supplied radio frequency radiation can provide sufficient power to apply a potential between a working electrode and a reference electrode sufficient to induce electrochemical reactions at the working electrode, measure the resulting amperometric current, and modulate the antenna impedance to adjust the backscatter radiation in a manner indicative of the measured amperometric current. In such an example, the supplied radio frequency radiation **171** can be considered an interrogation signal from the external reader **180** to the eye-mountable device **110** to request a measurement. By periodically interrogating the eye-mountable device **110** (e.g., by supplying radio frequency radiation **171** to temporarily turn the device on) and storing the sensor results (e.g., via the data storage **183**), the external reader **180** can accumulate a set of analyte concentration measurements over time without continuously powering the eye-mountable device **110**.

[0056] FIG. **2**A is a bottom view of an example eye-mountable electronic device **210**. FIG. **2**B is an aspect view of the example eye-mountable electronic device shown in FIG. **2**A. It is noted that relative dimensions in FIGS. **2**A and **2**B are not necessarily to scale, but have been rendered for purposes of explanation only in describing the arrangement of the example eye-mountable electronic device **210**. The eye-mountable device **210** is formed of a polymeric material **220** shaped as a curved disk. The polymeric material **220** can be a substantially transparent material to allow incident light to be transmitted to the eye while the eye-mountable device **210** is mounted to the eye. The polymeric material **220** can be a biocompatible material similar to those employed to form vision correction and/or cosmetic contact lenses in optometry, such as polyethylene terephthalate ("PET"), polymethyl methacrylate ("PMMA"), silicone hydrogels, polyhydroxyethylmethacrylate (polyHEMA) based hydrogels, and combinations of these, etc. The polymeric material **220** can be formed with one side having a concave surface **226** suitable to fit over a corneal surface of an eye. The opposing side of the disk can have a convex surface **224** that does not interfere with eyelid motion while the eye-mountable device **210** is mounted to the eye. A circular outer side edge **228** connects the concave surface **224** and convex surface **226**.

[0057] The eye-mountable device **210** can have dimensions similar to a vision correction and/or cosmetic contact lenses, such as a diameter of approximately 1 centimeter, and a thickness of about 0.1 to about 0.5 millimeters. However, the diameter and thickness values are provided for explanatory purposes only. In some embodiments, the dimensions of the eye-mountable device **210** can be selected according to the size and/or shape of the corneal surface of the wearer's eye.

[0058] The polymeric material **220** can be formed with a curved shape in a variety of ways. For example, techniques similar to those employed to form vision-correction contact lenses, such as heat molding, injection molding, spin casting, etc. can be employed to form the polymeric material **220**. While the eye-mountable device **210** is mounted in an eye, the convex surface **224** faces outward to the ambient environment while the concave surface **226** faces inward, toward the corneal surface. The convex surface **224** can therefore be considered an outer, top surface of the eye-mountable device **210** whereas the concave surface **226** can be considered an inner, bottom surface. The "bottom" view shown in FIG. **2**A is facing the concave surface **226**. From the bottom view shown in FIG. **2**A, the outer periphery **222**, near the outer circumference of the curved disk is curved out of the page, whereas the center region **221**, near the center of the disk is curved in to the page.

[0059] A substrate **230** is embedded in the polymeric material **220**. The substrate **230** can be embedded to be situated along the outer periphery **222** of the polymeric material **220**, away from the center region **221**. The substrate **230** does not interfere with vision because it is too close to the eye to be in focus and is positioned away from the center region **221** where incident light is transmitted to the eye-sensing portions of the eye. Moreover, the substrate **230** can be formed of a transparent material to further mitigate any effects on visual perception.

[0060] The substrate **230** can be shaped as a flat, circular ring (e.g., a disk with a central hole). The flat surface of the